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RECENTLY PUBLISHED RESEARCH OF INSTITUTE OF  
PHYSICAL CHEMISTRY IMENI L. V. PISARZHEVSKIY,  
ACADEMY OF SCIENCES USSR, MOSCOW

"Kinetics and Mechanism of Catalytic Conversion of Carbon Monoxide," V. A. Boyter, S. S. Gaukhman, N. P. Pisarzhevskaya, T. M. Gvaliya, Inst Phys Chem imeni L. V. Pisarzhevskiy, Acad Sci USSR, Moscow

"Zhur Priklad Khimii" Vol 18, 1945, pp 439-49

The dependence of yield of CO conversion to H and CO<sub>2</sub> in the presence of H<sub>2</sub>O was studied with the following factors: space velocity, total and partial pressures of the components, between 400° and 500° over K-carbon catalyst and Fe oxide catalyst of "nitrogen" type. The results are explained by a new theory for the K-carbon catalyst in which the following scheme of reactions is proposed:  $2\text{KOH} + \text{CO} \rightarrow \text{K}_2\text{CO}_3 + \text{H}_2$ ;  $\text{K}_2\text{CO}_3 + \text{H}_2 \rightarrow \text{K}_2\text{CO}_3 + \text{H}_2\text{O} \rightarrow 2\text{KOH} + \text{CO}_2$ , with summary result:  $\text{CO} + \text{H}_2\text{O} \rightarrow \text{H}_2 + \text{CO}_2$ . In case of the Fe catalysts there is evidence for considerable effect of the following sequence:  $2\text{CO} \rightarrow \text{CO}_2 + \text{C}$ ;  $\text{C} + 2\text{H}_2\text{O} \rightarrow \text{CO}_2 + 2\text{H}_2$ .

"Purification of Air by Removal of Admixture of Acetylene by Oxidation," V. A. Boyter, M. Ya. Rubanik, Yu. A. Snigurovskaya, E. A. Sadovnikova, Inst Phys Chem imeni L. V. Pisarzhevskiy, Acad Sci USSR, Moscow

"Zhur Priklad Khimii" Vol 18, 1945, pp 450-6

It was shown that  $C_2H_2$  can be removed from air by oxidation in passage over the following catalysts:

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MnO<sub>2</sub>, Mn ore promoted by Ag salts. The former is effective over a wide concentration range, the latter only for small concentrations. Various factors involved were studied. Generally, 1600° is a sufficiently high temperature for complete removal of C<sub>2</sub>H<sub>2</sub>.

"The Active Smooth Iron Electrode" B. Kabanov, D. Leikis, Inst Phys Chem imeni L. V. Pisarzhevskiy, Acad Sci USSR, Moscow

"Zhur Fiz Khimii" Vol 20, 1946, pp 995-1003

The rapid depolarization that distorts measurements of smooth Fe electrodes is avoided if the volume of the solution around the electrode is very small (e.g., 0.02 cc for an electrode surface of 3.5 sq cm). If such an electrode, after being heat-treated in H<sub>2</sub> at 7000 - 9000°, is immediately immersed in 2 N NaOH and anodically polarized with the current kept constant, its potential shows two breaks; e.g., for cd:  $2.4 \times 10^{-5}$  amp/sq cm these breaks are at +0.04 and +0.28 v (referred to H electrode) in 2 N NaOH. Extrapolation to zero current density yields the potentials -0.045 and +0.13 v. These presumably correspond to Fe | Fe(OH)<sub>2</sub> and to Fe | Fe(OH)<sub>2</sub>:Fe(OH)<sub>3</sub> electrodes. The amount of electricity consumed by an electrode along a break (i.e., the electrode capacity) is, e.g., 0.15-0.20 coulomb/sq cm of the visible surface area. Fe electrodes which had contact with the air before being placed in 2 N NaOH show no breaks. Their potential is increased from 0 to 0.5 v by as little as 0.0001 coulomb/sq cm. Presumably, this passivity of air-oxidized Fe is of the same type as the passivity of Pt.

"The Kinetics of Exothermal Catalytic Reactions in a Current: II. Theory of Reaction on a Short Contact Layer," Ya. L. Margolis, O. M. Todes, Inst Phys Chem imeni L. V. Pisarzhevskiy, Acad Sci USSR, Moscow

"Acta Physicochimica URSS" Vol 21, 1946, pp 88-98

Theoretical mathematical Equations are derived for a comparatively short reactor at not too small rates of flow. Two steady regimes, "quiet reactions" and "combustion regime," are established as in the previous case. For an unsteady extinction regime, the rate of cooling of the reactor and the activity of the catalyst can be connected by an almost linear relation; this permits the development of a simple method of evaluating the relative activities of catalysts for a given reaction.

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"Poisoning and Modifying Catalysts," S. Z. Roginskiy,  
Inst Phys Chem imeni L. V. Pisarzhevskiy, Acad Sci  
USSR, Moscow

"Zhur Fiz Khimii" Vol 21, 1947, pp 1143-58

A review of the earlier experimental work shows that poisoning and promoting of catalysts by additions are not as different as believed hitherto. The mechanism usually postulated for poisoning and promoting cannot account for the observations. A definite impurity can either raise or lower the efficiency of a catalyst according to its amount. The effect of a definite amount of an impurity can be positive or negative according to conditions. The increase of efficiency is not always due to a reduction in the activation. Catalysts can be modified by additions, not only poisoned or promoted.

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